## AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application:

## LISTING OF CLAIMS:

- 1. (currently amended) A transgenic non-human animal mouse expressing at least one transgene comprising a DNA sequence encoding a heterologous Amyloid Precursor Protein(APP) comprising at least the Arctic mutation (E693G) and a further Alzeihmer's disease (AD) AD (Alzheimer's disease) pathogenic mutation or a further transgene affecting AD pathogenesis, which results in increased amounts of intracellular soluble A $\beta$  aggregates, including A $\beta$  peptides.
- 2. (currently amended) The transgenic animal mouse according to claim 1, wherein the transgene/transgenes are integrated in the genomic DNA.
- 3. (currently amended) The transgenic animal mouse according to claim 1, wherein said transgene/transgenes are operably linked to a promoter effective for expression of said gene in the brain tissue of said animal mouse.

4. (currently amended) The transgenic  $\frac{\text{animal }}{\text{mouse}}$  according to claim 1, wherein the endogenous APP is expressive or non-expressive.

## 5. (cancelled)

- 6. (withdrawn-currently amended) The transgenic animal  $\underline{\text{mouse}}$  according to claim 1, wherein said further transgene comprises a DNA sequence encoding apolipoprotein E, apolipoprotein J(clusterin),  $a_1$ -antichymotrypsin (ACT) or fragments thereof.
- 7. (currently amended) The transgenic animal mouse according to claim 1, wherein said further AD pathogenic mutation is one of the APP mutations KM670/671DF, KM670/671DY, KM670/671EF or KM670/671EY.
- 8. (currently amended) The transgenic animal mouse according to claim 1, wherein said further AD pathogenic mutation is one of the APP mutations KM670/671NL, KM670/671NY, KM670/671NF, KM670/671KL, KM670/671DL or KM670/671EL, wherein KM670/671NL (the Swedish mutation) is preferred.
- 9. (currently amended) The transgenic animal mouse according to claim 1, wherein the transgenic animal mouse

expresses only one transgene which comprises only  $\underline{\text{E693G}}$  and  $\underline{\text{KM670/671NL}}$  the Arctic mutation ( $\underline{\text{E693G}}$ ) and the Swedish mutation ( $\underline{\text{KM670/671NL}}$ ).

10. (withdrawn-currently amended) The transgenic animal mouse according to claim 1, additionally comprising a homologously integrated targeting construct for at least one of the neprilysin or insulin-degrading enzyme (IDE) genes, which disrupts these genes through gene ablation (knock-out) and enhances  $A\beta-40$  and/or  $A\beta-42$  Arctic peptide production.

# 11-13. (cancelled)

14. (currently amended) A method of producing the transgenic animal mouse according to claim 1, comprising:

by microinjecting said at least one transgene into a fertilized egg or an embryo, said at least one transgene comprising a DNA sequence encoding a heterologous Amyloid Precursor Protein (APP) comprising at least the Arctic mutation (E693G) and a further AD (Alzheimer's disease) Alzeihmer's disease (AD) pathogenic mutation or a further transgene affecting AD pathogenesis;

transferring said fertilized egg or said embryo microinjected with said at least one transgene to a mouse so as

to produce a transgenic mouse from said fertilized egg or said embryo.

- 15. (currently amended) The method according to claim 14, wherein said transgene/transgenes are operably linked to a promoter effective for expression of said gene in the brain tissue of said  $\frac{1}{2}$  mouse.
- 16. (withdrawn) The method according to claim 14, wherein the endogenous APP is optionally made non-expressive.

#### 17. (cancelled)

- 18. (withdrawn) The method according to claim 14, wherein said further transgene comprises a DNA sequence encoding apolipoprotein E, apolipoprotein J (clusterin), al-antichymotrypsin (ACT) or fragments thereof.
- 19. (previously presented) The method according to claim 14, wherein said further AD pathogenic mutation is one of the APP mutations KM670/671DF, KM670/671DY, KM670/671EF or KM670/671EY.
- 20. (currently amended) The method according to claim 14, wherein said further AD pathogenic mutation is one of

the APP mutations KM670/671NL, KM670/671NY, KM670/671NF, KM670/671KL, KM670/671DL or KM670/671EL, wherein KM670/671NL (the Swedish mutation) is preferred.

- 21. (withdrawn) The method according to claim 14, additionally comprising homologously integrating a targeting construct for at least one of the neprilysin or insulin-degrading enzyme (IDE) genes.
- 22. (currently amended) A method of screening <u>agents</u> useful for treating, preventing or inhibiting Alzheimer's <u>disease</u>, <u>comprising</u>:

administering wherein the an agent to a first transgenic animal mouse according to claim 1 is used for screening for agents; and

observing the ability of the first transgenic mouse to form  ${\tt A}{\beta}$  peptides;

comparing the ability of the first transgenic mouse to form  $A\beta$  peptides to the ability of a second transgenic mouse according to claim 1 to form  $A\beta$  peptides, the agent not being administered to the second transgenic mouse;

 $\frac{\text{wherein a decrease in } A\beta \text{ formation in the first}}{\text{transgenic mouse indicates that the agent is}} \text{ useful for treating,}$  preventing or inhibiting Alzheimer's disease.

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23. (currently amended) A method of screening <u>for</u> diagnostic agents for Alzheimer's , <u>comprising:</u>

administering an agent to a first transgenic mouse according to claim 1 an agent;

comparing the ability of the first transgenic mouse to form  $A\beta$  peptides to the ability of a second transgenic mouse according to claim 1 to form  $A\beta$  peptides, the agent not being administered to the second transgenic mouse;

wherein a decrease in A $\beta$  formation in the first transgenic mouse indicates that the agent wherein the transgenic animal according to claim 1 is used for screening for diagnostic agents is a diagnostic agent for Alzheimer's disease.

- 24. (new) The transgenic mouse according to claim 8, wherein said further AD pathogenic mutation is KM670/671NL.
- 25. (new) The method according to claim 20, wherein said further AD pathogenic mutation is KM670/671NL.